Opis choroby *

Definicja

A rare glomerular disease, histologically characterized by thickening of the capillary wall, with immune deposits predominantly containing IgG4 and C3 on the sub-epithelial side, and typically manifesting with nephrotic syndrome.

Dane

Klasyfikacja
Choroba

Synonimy
Idiopathic membranous glomerulonephritis
Primary membranous nephropathy
Idiopathic membranous glomerulonephritis
Primary membranous nephropathy

Kod ORPHA
97560

Kod OMIM
614692

Kod ICD10
N04.2

Kod ICD11
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*Źródło

orphanet